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(54) Title: LASER MARKING METHOD

(57) Abstract: A method of colouring a polymeric material containing a) a latent acid, b) a colour former and c) optionally further ingredients by irradiation with UV-light.

Laser marking method

The present application relates to a method of coloring a polymeric material on irradiation with a laser emitting UV-light.

Recently, for real-time marking of letters and signs such as marker's name, product name, date of production, lot number etc. on the surfaces of various commercial articles, the laser marking system is popularly employed for its various advantages. However, the existing laser marking systems do not perfectly fulfill all the user's requirements and thus a need exists to improve the properties of such systems.

It has now been found that the inventive laser marking method gives excellent results and is applicable on various materials.

The present invention relates to a method of coloring a polymeric material, wherein a polymeric material containing

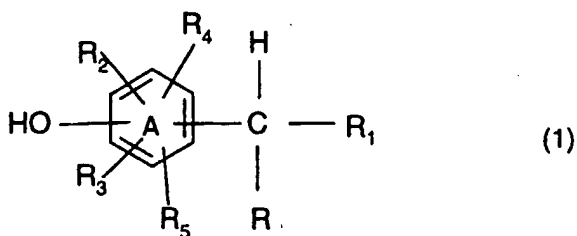
- a) a latent acid,
 - b) a colour former and
 - c) optionally further ingredients
- is irradiated with UV-light.

If a laser emits the UV-light, usually a clear cut mark is obtained. A clear cut mark means that the polymeric material is coloured at only those areas which have been irradiated with UV-light. The latent acid absorbs the energy of the UV-light and is transformed into the corresponding acid, which in turn reacts with the colour former, which itself turns into a coloured material.

As latent acids compounds are suitable which are not acids per se and contain a proton, which can be split off by irradiation.

Preferred latent acids are of formula

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wherein ring A can contain one or more hetero atoms and/or can contain an anelated ring, R_1 is hydrogen, alkyl, preferably C_1 - C_{20} -alkyl, alkenyl, preferably C_2 - C_{20} -alkenyl, aryl, preferably phenyl or phenyl which is substituted one to three times with C_1 - C_4 alkyl, or C_1 - C_4 alkoxy,

R_2 , R_3 , R_4 and R_5 independently of each other are hydrogen or a functional substituent, and R stands for C_1 - C_6 alkyl, $-Z_1-Q_1$, or $-Z_2-Q_2$,

wherein Z_1 is a single bond, S, NH or O, and Q_1 is a heterocyclic ring system having from 5 to 9 ring atoms selected from C, S, O and N, with at least 2, preferably at least 3, more preferably at least 4 carbon atoms in the ring system, preferably Q_1 stands for morpholine, pyridine, which may be substituted one to three times with C_1 - C_4 alkyl or hydroxy, mercaptobenzoxazole, mercaptobenzthiazole,

and wherein Z_2 stands for C_1 - C_4 alkylene, which can be substituted by C_1 - C_4 alkyl or Q_3 , wherein Q_3 stands for phenyl which can be substituted one to three times with C_1 - C_4 alkyl, hydroxy, C_5 - C_8 cycloalkyl and/or a heterocyclic ring system having from 5 to 9 ring atoms selected from C, S, O and N, with at least 2, preferably at least 3, more preferably at least 4 carbon atoms in the ring system, and Q_2 stands for phenyl which can be substituted one to three times with C_1 - C_4 alkyl, hydroxy, C_5 - C_8 cycloalkyl and/or a heterocyclic ring system having from 5 to 9 ring atoms selected from C, S, O and N, with at least 2, preferably at least 3, more preferably at least 4 carbon atoms in the ring system, with the proviso that the hydrogen atom at the C-atom in α -position to R can be split off by irradiation.

Suitable rings A are e.g. phenyl, naphthyl, pyridyl and quinoliny, phenyl and pyridyl are especially preferred.

Preferably, Z_2 stands for $-CH_2-$, $-CH_2-CH_2-$, $-CH_2-CHMe-$, $-CH_2-CHQ_3-$, in which Q_3 stands for 4-hydroxy-3-i-propyl-6-methylphenyl, 4-hydroxy-3-tert.-butyl-6-methylphenyl, or 4-hydroxy-3-cyclohexyl-6-methylphenyl and Q_2 stands for phenyl or 4-hydroxy-3-i-propyl-6-

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methylphenyl, 4-hydroxy-3-tert.-butyl-6-methylphenyl, or 4-hydroxy-3-cyclohexyl-6-methylphenyl.

R₁ is preferably hydrogen, or methyl.

Functional substituents R₂, R₃, R₄ and R₅ are e. g. C₁-C₂₀-alkyl, preferably C₁-C₈-alkyl, particularly preferred C₁-C₆-alkyl, especially preferred C₁-C₄-alkyl, C₅-C₈-cycloalkyl, C₂-C₂₀-alkenyl, preferred C₂-C₆-alkenyl, C₁-C₆-alkoxy, hydroxy, halogen, nitro, cyano, -SO₂R', wherein R' is hydrogen, alkyl or a metallic cation such as a alkali metal, e.g. sodium or potassium, or earth alkali metal cation, e.g. calcium, or phenyl, which may be substituted one to three times with hydroxy and/or Z₂₁-R₇, wherein Z₂₁ stands for C₁-C₄alkylene, which can be substituted by C₁-C₄alkyl, and R₇ stands for hydrogen, C₁-C₄alkyl or phenyl, which may be substituted one to three times with hydroxy, C₁-C₄alkyl and/or Z₂₂-R₈, wherein Z₂₂ stands for C₁-C₄alkylene, which can be substituted by C₁-C₄alkyl, and R₈ stands for a heterocyclic ring system having from 5 to 9 ring atoms selected from C, S, O and N, with at least 2, preferably at least 3, more preferably at least 4 carbon atoms in the ring system, preferably R₈ stands for morpholine. In a preferred embodiment of this invention R₂, R₃, R₄ and R₅ are preferably independently of each other hydrogen, C₁-C₂₀-alkyl or C₂-C₂₀-alkenyl or substituted phenyl wherein hydroxy and Z₂₁-R₇ being the substituents. Especially preferred compounds of formula (1) are those wherein R₂ and R₃ are independently of each other C₁-C₈-alkyl and R₄ and R₅ are each hydrogen.

Halogen means fluoro, chloro, bromo, or iodo, preferably chloro.

Heterocyclic residue or heterocyclic ring system having at least 2, preferably at least 3, more preferably at least 4 carbon atoms means e.g. an optionally substituted monocyclic or bicyclic heterocyclic residue such as pyrrolidino, piperidino, morpholino, benzthiazole, 1,2,4-triazole, imidazole, pyrazole, tetrazole, thiazolin-2-thione, imidazolin-2-thione, N-methyl-imidazolin-2-thione and 5-(3-phenyl-1,3,4-thia-diazol-2(3H)-thione), 2-pyridine, 4-pyridine, 3-pyridazine, 2-pyrimidine, 2-thiazole, 2-thioazoline, 3-(1,2,4-triazole) and 5-(2-mercapto-1,3,4-thiadiazole), naphthyridine, purine and pteridine residues, benzimidazole, benzotriazole, benzoxazolin-2-thione, 2-benzoxazole, mercaptobenzoxazol, mercaptobenzthiazol and quinolinyl.

It is furthermore preferred that at least one of R_2 and R_3 is in o-position to the OH-group.

The organic residues R and R_1 can be independently of each other of any kind with the proviso that the H at the C-atom in α -position to R can be split off by irradiation.

Preferably R is a heterocyclic residue which is bond to the CHR_1 -group via a nitrogen, oxygen or sulfur atom or is a C_1 - C_6 -alkyl which is unsubstituted or substituted, e.g. by hydroxy, C_1 - C_6 -alkoxy or unsubstituted or substituted aryl, especially phenyl. Suitable substituents for aryl are preferably the above mentioned substituents R_2 through R_5 . Most preferably R is a radical of mercaptobenzoxazol or mercaptobenzthiazol or C_1 - C_4 -alkyl which is unsubstituted or substituted by unsubstituted phenyl or phenyl carrying 1 to 4 substituents selected from the group consisting of C_1 - C_6 -alkyl, C_1 - C_4 -alkox and hydroxy.

In preferred compounds of formula (1) the residue $-CHRR_1$ is situated in o- or especially in p-position to the OH-group.

C_1 - C_{20} -alkyl means e.g. methyl, ethyl, n-, i-propyl, n-, sec-, iso-, tert.-butyl, n-pentyl, n-hexyl, n-heptyl, n-octyl, n-nonyl, n-decyl, n-undecyl, n-dodecyl, n-tridecyl, n-tetradecyl, n-pentadecyl, n-hexadecyl, n-heptadecyl, n-octadecyl, n-nonadecyl, n-eicosyl, preferably C_1 - C_6 -alkyl such as methyl, ethyl, n-, i-propyl, n-, sec-, iso-, tert.-butyl, n-pentyl, n-hexyl, n-heptyl, n-octyl, particularly preferred C_1 - C_6 -alkyl such as methyl, ethyl, n-, i-propyl, n-, sec-, iso-, tert.-butyl, n-pentyl, n-hexyl, especially preferred C_1 - C_4 -alkyl such as methyl, ethyl, n-, i-propyl, n-, sec-, iso-, tert.-butyl.

C_5 - C_8 -cycloalkyl stands for cyclopentyl, cyclohexyl, cycloheptyl, or cyclooctyl, preferably cyclohexyl.

C_2 - C_{20} -alkenyl stands for e.g. ethenyl, n-, i-propenyl, n-, sec-, iso-, tert.-butenyl, n-pentenyl, n-hexenyl, n-heptenyl, n-octenyl, n-nonenyl, n-decenyl, n-undecenyl, n-dodecenyl, n-tridecenyl, n-tetradecenyl, n-pentadecenyl, n-hexadecenyl, n-heptadecenyl, n-octadecenyl, n-nonadecenyl, n-eicosenyl, preferably C_2 - C_6 -alkyl such as ethenyl, n-, i-propenyl, n-, sec-, iso-, tert.-butenyl, n-pentenyl, n-hexenyl.

C₁-C₈-alkoxy stands for e.g. methoxy, ethoxy, n-, i-propoxy, n-, sec-, iso-, tert.-butoxy, n-pentoxy, n-hexoxy.

Preferred polymeric material according to the present invention contains a latent acid of formula (1) wherein

the ring A is phenyl or pyridyl,

R₁ is hydrogen,

R₂ and R₃ are independently of each other C₁-C₄-alkyl,

R₄ and R₅ are each hydrogen and

R is a heterocyclic residue which is bond to the CHR₁-group via a nitrogen, oxygen or sulfur atom or is a C₁-C₈-alkyl which is unsubstituted or substituted.

Preferred polymeric material according to the present invention contains a latent acid of formula (1) wherein

the ring A is phenyl or pyridyl,

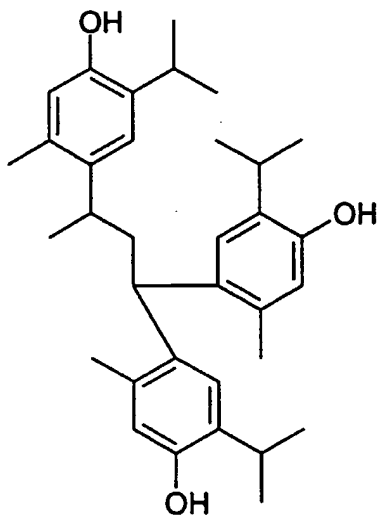
R₁ is hydrogen.

R₂ and R₃ are independently of each other C₁-C₄-alkyl,

R₄ and R₅ are each hydrogen and

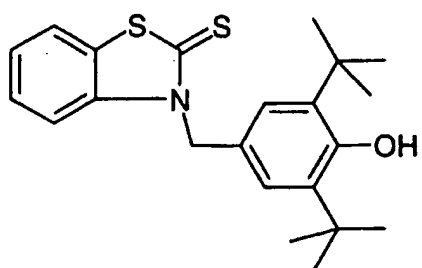
R is a heterocyclic residue, which is bond via a nitrogen, oxygen or sulfur atom or is a C₁-C₈-alkyl, which is unsubstituted or substituted.

Especially preferred compounds of formula (1) are the following compounds:

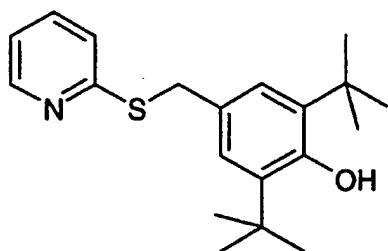


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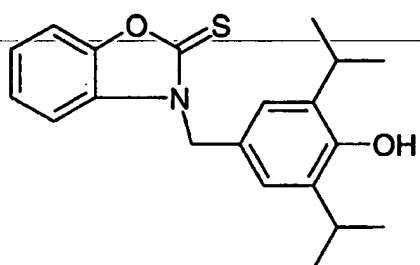
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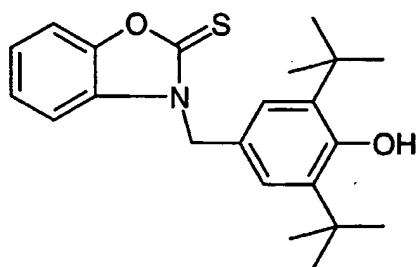
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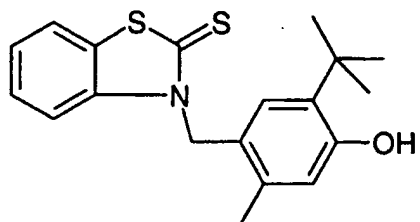
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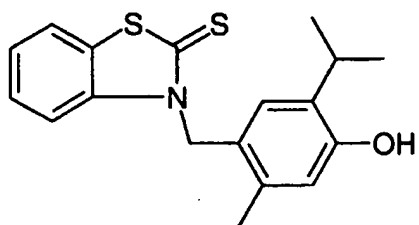
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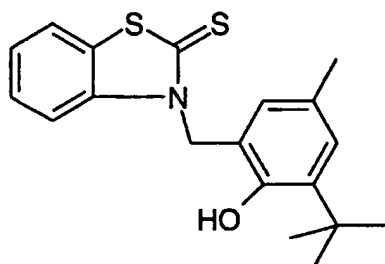
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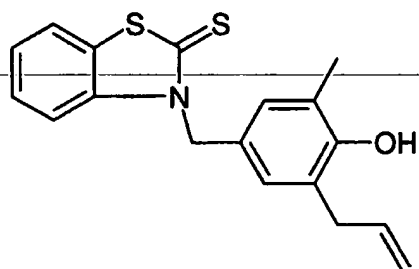
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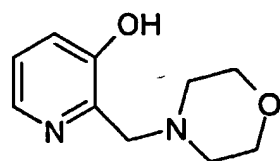
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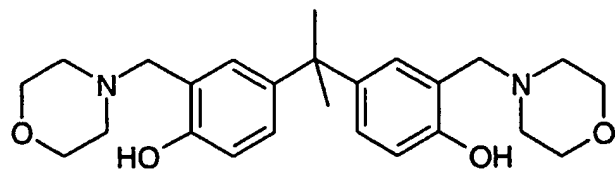
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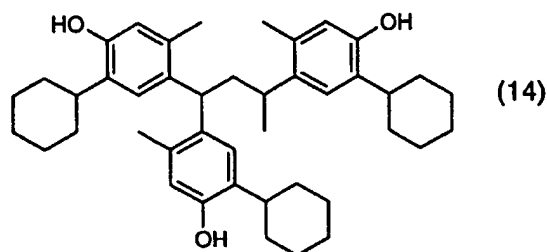
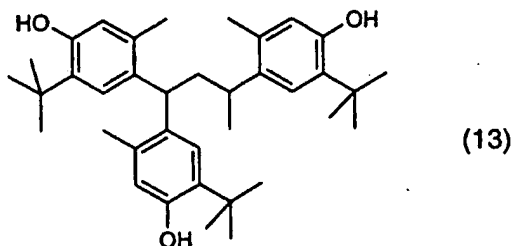


(11)



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The compounds of formula (1) are known or can be made in a manner known per se, e. g. the compound (2) according to GB 2,120,243 and compounds (7) and (8) as described in EP-A-330 613.

Polymeric material useable for the present invention is preferably synthetic organic polymeric material, especially material commonly used for electronic applications.

In particular the following polymers are preferred:

1. Polymers of monoolefins and diolefins, for example polypropylene, polyisobutylene, polybut-1-ene, poly-4-methylpent-1-ene, polyvinylcyclohexane, polyisoprene or polybutadiene, as well as polymers of cycloolefins, for instance of cyclopentene or norbornene, polyethylene (which optionally can be crosslinked), for example high density polyethylene (HDPE), high density and high molecular weight polyethylene (HDPE-HMW), high density and ultrahigh molecular weight polyethylene (HDPE-UHMW), medium density polyethylene (MDPE), low density polyethylene (LDPE), linear low density polyethylene (LLDPE), (VLDPE) and (ULDPE).

Polyolefins, i.e. the polymers of monoolefins exemplified in the preceding paragraph, preferably polyethylene and polypropylene, can be prepared by different, and especially by the following, methods:

a) radical polymerisation (normally under high pressure and at elevated temperature).

- b) catalytic polymerisation using a catalyst that normally contains one or more than one metal of groups IVb, Vb, VIb or VIII of the Periodic Table. These metals usually have one or more than one ligand, typically oxides, halides, alcoholates, esters, ethers, amines, alkyls, alkenyls and/or aryls that may be either π - or σ -coordinated. These metal complexes may be in the free form or fixed on substrates, typically on activated magnesium chloride, titanium(III) chloride, alumina or silicon oxide. These catalysts may be soluble or insoluble in the polymerisation medium. The catalysts can be used by themselves in the polymerisation or further activators may be used, typically metal alkyls, metal hydrides, metal alkyl halides, metal alkyl oxides or metal alkyloxanes, said metals being elements of groups Ia, IIa and/or IIIa of the Periodic Table. The activators may be modified conveniently with further ester, ether, amine or silyl ether groups. These catalyst systems are usually termed Phillips, Standard Oil Indiana, Ziegler (-Natta), TNZ (DuPont), metallocene or single site catalysts (SSC).
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2. Mixtures of the polymers mentioned under 1), for example mixtures of polypropylene with polyisobutylene, polypropylene with polyethylene (for example PP/HDPE, PP/LDPE) and mixtures of different types of polyethylene (for example LDPE/HDPE).

3. Copolymers of monoolefins and diolefins with each other or with other vinyl monomers, for example ethylene/propylene copolymers, linear low density polyethylene (LLDPE) and mixtures thereof with low density polyethylene (LDPE), propylene/but-1-ene copolymers, propylene/isobutylene copolymers, ethylene/but-1-ene copolymers, ethylene/hexene copolymers, ethylene/methylpentene copolymers, ethylene/heptene copolymers, ethylene/octene copolymers, ethylene/vinylcyclohexane copolymers, ethylene/cycloolefin copolymers (e.g. ethylene/norbornene like COC), ethylene/1-olefins copolymers, where the 1-olefin is generated in-situ; propylene/butadiene copolymers, isobutylene/isoprene copolymers, ethylene/vinylcyclohexene copolymers, ethylene/alkyl acrylate copolymers, ethylene/alkyl methacrylate copolymers, ethylene/vinyl acetate copolymers or ethylene/acrylic acid copolymers and their salts (ionomers) as well as terpolymers of ethylene with propylene and a diene such as hexadiene, dicyclopentadiene or ethylidene-norbornene; and mixtures of such copolymers with one another and with polymers mentioned in 1) above, for example polypropylene/ethylene-propylene copolymers, LDPE/ethylene-vinyl acetate copolymers (EVA), LDPE/ethylene-

acrylic acid copolymers (EAA), LLDPE/EVA, LLDPE/EAA and alternating or random polyalkylene/carbon monoxide copolymers and mixtures thereof with other polymers, for example polyamides.

4. Hydrocarbon resins (for example C₅-C₉) including hydrogenated modifications thereof (e.g. tackifiers) and mixtures of polyalkylenes and starch.

Homopolymers and copolymers from 1.) - 4.) may have any stereostructure including syndiotactic, isotactic, hemi-isotactic or atactic; where atactic polymers are preferred. Stereoblock polymers are also included.

5. Polystyrene, poly(p-methylstyrene), poly(α -methylstyrene).

6. Aromatic homopolymers and copolymers derived from vinyl aromatic monomers including styrene, α -methylstyrene, all isomers of vinyl toluene, especially p-vinyltoluene, all isomers of ethyl styrene, propyl styrene, vinyl biphenyl, vinyl naphthalene, and vinyl anthracene, and mixtures thereof. Homopolymers and copolymers may have any stereostructure including syndiotactic, isotactic, hemi-isotactic or atactic; where atactic polymers are preferred. Stereoblock polymers are also included.

6a. Copolymers including aforementioned vinyl aromatic monomers and comonomers selected from ethylene, propylene, dienes, nitriles, acids, maleic anhydrides, maleimides, vinyl acetate and vinyl chloride or acrylic derivatives and mixtures thereof, for example styrene/butadiene, styrene/acrylonitrile, styrene/ethylene (interpolymers), styrene/alkyl methacrylate, styrene/butadiene/alkyl acrylate, styrene/butadiene/alkyl methacrylate, styrene/maleic anhydride, styrene/acrylonitrile/methyl acrylate; mixtures of high impact strength of styrene copolymers and another polymer, for example a polyacrylate, a diene polymer or an ethylene/propylene/diene terpolymer; and block copolymers of styrene such as styrene/butadiene/styrene, styrene/isoprene/styrene, styrene/ethylene/butylene/styrene or styrene/ethylene/propylene/styrene.

6b. Hydrogenated aromatic polymers derived from hydrogenation of polymers mentioned under 6.), especially including polycyclohexylethylene (PCHE) prepared by hydrogenating atactic polystyrene, often referred to as polyvinylcyclohexane (PVCH).

6c. Hydrogenated aromatic polymers derived from hydrogenation of polymers mentioned under 6a.).

Homopolymers and copolymers may have any stereostructure including syndiotactic, isotactic, hemi-isotactic or atactic; where atactic polymers are preferred. Stereoblock polymers are also included.

7. Graft copolymers of vinyl aromatic monomers such as styrene or α -methylstyrene, for example styrene on polybutadiene, styrene on polybutadiene-styrene or polybutadiene-acrylonitrile copolymers; styrene and acrylonitrile (or methacrylonitrile) on polybutadiene; styrene, acrylonitrile and methyl methacrylate on polybutadiene; styrene and maleic anhydride on polybutadiene; styrene, acrylonitrile and maleic anhydride or maleimide on polybutadiene; styrene and maleimide on polybutadiene; styrene and alkyl acrylates or methacrylates on polybutadiene; styrene and acrylonitrile on ethylene/propylene/diene terpolymers; styrene and acrylonitrile on polyalkyl acrylates or polyalkyl methacrylates, styrene and acrylonitrile on acrylate/butadiene copolymers, as well as mixtures thereof with the copolymers listed under 6), for example the copolymer mixtures known as ABS, MBS, ASA or AES polymers.

8. Halogen-containing polymers such as polychloroprene, chlorinated rubbers, chlorinated and brominated copolymer of isobutylene-isoprene (halobutyl rubber), chlorinated or sulfochlorinated polyethylene, copolymers of ethylene and chlorinated ethylene, epichlorohydrin homo- and copolymers, especially polymers of halogen-containing vinyl compounds, for example polyvinyl chloride, polyvinylidene chloride, polyvinyl fluoride, polyvinylidene fluoride, as well as copolymers thereof such as vinyl chloride/vinylidene chloride, vinyl chloride/vinyl acetate or vinylidene chloride/vinyl acetate copolymers.

9. Polymers derived from α,β -unsaturated acids and derivatives thereof such as polyacrylates and polymethacrylates; polymethyl methacrylates, polyacrylamides and polyacrylonitriles, impact-modified with butyl acrylate.

10. Copolymers of the monomers mentioned under 9) with each other or with other unsaturated monomers, for example acrylonitrile/ butadiene copolymers, acrylonitrile/alkyl acrylate

copolymers, acrylonitrile/alkoxyalkyl acrylate or acrylonitrile/vinyl halide copolymers or acrylonitrile/ alkyl methacrylate/butadiene terpolymers.

11. Polymers derived from unsaturated alcohols and amines or the acyl derivatives or acetals thereof, for example polyvinyl alcohol, polyvinyl acetate, polyvinyl stearate, polyvinyl benzoate, polyvinyl maleate, polyvinyl butyral, polyallyl phthalate or polyallyl melamine; as well as their copolymers with olefins mentioned in 1) above.

12. Homopolymers and copolymers of cyclic ethers such as polyalkylene glycols, polyethylene oxide, polypropylene oxide or copolymers thereof with bisglycidyl ethers.

13. Polyacetals such as polyoxymethylene and those polyoxymethylenes, which contain ethylene oxide as a comonomer; polyacetals modified with thermoplastic polyurethanes, acrylates or MBS.

14. Polyphenylene oxides and sulfides, and mixtures of polyphenylene oxides with styrene polymers or polyamides.

15. Polyurethanes derived from hydroxyl-terminated polyethers, polyesters or polybutadienes on the one hand and aliphatic or aromatic polyisocyanates on the other, as well as precursors thereof.

16. Polyamides and copolyamides derived from diamines and dicarboxylic acids and/or from aminocarboxylic acids or the corresponding lactams, for example polyamide 4, polyamide 6, polyamide 6/6, 6/10, 6/9, 6/12, 4/6, 12/12, polyamide 11, polyamide 12, aromatic polyamides starting from m-xylene diamine and adipic acid; polyamides prepared from hexamethylenediamine and isophthalic or/and terephthalic acid and with or without an elastomer as modifier, for example poly-2,4,4-trimethylhexamethylene terephthalamide or poly-m-phenylene isophthalamide; and also block copolymers of the aforementioned polyamides with polyolefins, olefin copolymers, ionomers or chemically bonded or grafted elastomers; or with polyethers, e.g. with polyethylene glycol, polypropylene glycol or polytetramethylene glycol; as well as polyamides or copolyamides modified with EPDM or ABS; and polyamides condensed during processing (RIM polyamide systems).

17. Polyureas, polyimides, polyamide-imides, polyetherimids, polyesterimids, polyhydantoins and polybenzimidazoles.

18. Polyesters derived from dicarboxylic acids and diols and/or from hydroxycarboxylic acids or the corresponding lactones, for example polyethylene terephthalate, polybutylene terephthalate, poly-1,4-dimethylolcyclohexane terephthalate, polyalkylene naphthalate (PAN) and polyhydroxybenzoates, as well as block copolyether esters derived from hydroxyl-terminated polyethers; and also polyesters modified with polycarbonates or MBS.

19. Polycarbonates and polyester carbonates.

20. Polyketones.

21. Polysulfones, polyether sulfones and polyether ketones.

22. Crosslinked polymers derived from aldehydes on the one hand and phenols, ureas and melamines on the other hand, such as phenol/formaldehyde resins, urea/formaldehyde resins and melamine/formaldehyde resins.

23. Drying and non-drying alkyd resins.

24. Unsaturated polyester resins derived from copolyesters of saturated and unsaturated dicarboxylic acids with polyhydric alcohols and vinyl compounds as crosslinking agents, and also halogen-containing modifications thereof of low flammability.

25. Crosslinkable acrylic resins derived from substituted acrylates, for example epoxy acrylates, urethane acrylates or polyester acrylates.

26. Alkyd resins, polyester resins and acrylate resins crosslinked with melamine resins, urea resins, isocyanates, isocyanurates, polyisocyanates or epoxy resins.

27. Crosslinked epoxy resins derived from aliphatic, cycloaliphatic, heterocyclic or aromatic glycidyl compounds, e.g. products of diglycidyl ethers of bisphenol A and bisphenol F, which

are crosslinked with customary hardeners such as anhydrides or amines, with or without accelerators.

28. Natural polymers such as cellulose, rubber, gelatin and chemically modified homologous derivatives thereof, for example cellulose acetates, cellulose propionates and cellulose butyrates, or the cellulose ethers such as methyl cellulose; as well as rosins and their derivatives.

29. Blends of the aforementioned polymers (polyblends), for example PP/EPDM, Polyamide/EPDM or ABS, PVC/EVA, PVC/ABS, PVC/MBS, PC/ABS, PBTP/ABS, PC/ASA, PC/PBT, PVC/CPE, PVC/acrylates, POM/thermoplastic PUR, PC/thermoplastic PUR, POM/acrylate, POM/MBS, PPO/HIPS, PPO/PA 6.6 and copolymers, PA/HDPE, PA/PP, PA/PPO, PBT/PC/ABS or PBT/PET/PC.

Especially preferred is organic polymeric material made of SAN (copolymer made of styrene and acrylonitrile), PP (polypropylene), PE (polyethylene), PVC (polyvinylchloride), PET (polyethyleneterephthalate), PET-G (glycole-modified PET), PMMA (polymethylmethacrylate) and related polyacrylics, PS (polystyrene), ASA (copolymer made of acrylonitrile, styrene, acrylate), PA (polyamide), ABS (copolymer made of acrylonitrile, styrene, butadiene), LLDPE (linear LDPE), LDPE (low density polyethylene), HDPE (high density polyethylene) and polycarbonate, most preferably polycarbonate. The polymeric material can also be a mixture of two or more different polymers.

The colour forming compounds are, for example, triphenylmethanes, lactones, benzoxazines, spiropyrans or preferably fluorans or phthalides.

Suitable colour formers include but are not limited to; 3-dibutylamino-7-dibenzylaminofluoran, 3-diethylamino-6-methylfluoran, 3-dimethylamino-6-methyl-7-anilino-7-fluoran, 3-diethylamino-6-methyl-7-anilino-7-fluoran, 3-diethylamino-6-methyl-7-(2,4-dimethylanilino) fluoran, 3-diethylamino-6-methyl-7-chlorofluoran, 3-diethylamino-6-methyl-7-(3-trifluoromethylanilino) fluoran, 3-diethylamino-6-methyl-7-(2-chloroanilino) fluoran, 3-diethylamino-6-methyl-7-(4-chloroanilino) fluoran, 3-diethylamino-6-methyl-7-(2-fluoroanilino) fluoran, 3-diethylamino-6-methyl-7-(4-n-octylanilino) fluoran, 3-diethylamino-7-(4-n-octylanilino) fluoran, 3-diethylamino-7-(4-n-octylamino) fluoran, 3-diethylamino-6-methyl-7-(dibenzylamino) fluoran, 3-

diethylamino-7-(dibenzylamino) fluoran, 3-diethylamino-6-chloro-7-methylfluoran, 3-diethylamino-7-t-butylfluoran, 3-diethylamino-7-carboxyethylfluoran, 3-diethylamino-6-chloro-7-anilinofluoran, 3-diethylamino-6-methyl-7-(3-methylanilino) fluoran, 3-diethylamino-6-methyl-7-(4-methylanilino) fluoran, 3-diethylamino-6-ethoxyethyl-7-anilinofluoran, 3-diethylamino-7-methylfluoran, 3-diethylamino-7-chlorofluoran, 3-diethylamino-7-(3-trifluoromethylanilino) fluoran, 3-diethylamino-7-(2-chloroanilino) fluoran, 3-diethylamino-7-(2-fluoroanilino) fluoran, 3-diethylamino-benzo[a] fluoran, 3-diethylamino-benzo[c] fluoran, 3-dibutylamino-6-methyl fluoran, 3-dibutylamino-6-methyl-7-anilinofluoran, 3-dibutylamino-6-methyl-7-(2,4-dimethylanilino) fluoran, 3-dibutylamino-6-methyl-7-(2-chloroanilino) fluoran, 3-dibutylamino-6-methyl-7-(4-chloroanilino) fluoran, 3-dibutylamino-6-methyl-7-(2-fluoroanilino) fluoran, 3-dibutylamino-6-methyl-7-(3-trifluoromethylanilino) fluoran, 3-dibutylamino-6-ethoxyethyl-7-anilinofluoran, 3-dibutylamino-6-chloro-anilinofluoran, 3-dibutylamino-6-methyl-7-(4-methylanilino) fluoran, 3-dibutylamino-7-(2-chloroanilino) fluoran, 3-dibutylamino-7-(2-fluoroanilino) fluoran, 3-dipentylamino-6-methyl-7-anilinofluoran, 3-dipentylamino-6-methyl-7-(4-2-chloroanilino) fluoran, 3-dipentylamino-7-(3-trifluoromethylanilino) fluoran, 3-dipentylamino-6-chloro-7-anilinofluoran, 3-dipentylamino-7-(4-chloroanilino) fluoran, 3-pyrrolidino-6-methyl-7-anilinofluoran, 3-piperidino-6-methyl-7-anilinofluoran, 3-(N-methyl-N-propylamino)-6-methyl-7-anilinofluoran, 3-(N-methyl-N-cyclohexylamino)-6-methyl-7-anilinofluoran, 3-(N-ethyl-N-cyclohexylamino)-6-methyl-7-anilinofluoran, 3-(N-ethyl-p-toluidino)-6-methyl-7-anilinofluoran, 3-(N-ethyl-N-isoamylamino)-6-methyl-7-anilinofluoran, 3-(N-ethyl-N-isoamylamino)-6-chloro-7-anilinofluoran, 3-(N-ethyl-N-tetrahydrofurfurylamino)-6-methyl-7-anilinofluoran, 3-(N-ethyl-N-isobutylamino)-6-methyl-7-anilinofluoran, 3-(N-butyl-N-isoamylamino)-6-methyl-7-anilinofluoran, 3-(N-isopropyl-N-3-pentylamino)-6-methyl-7-anilinofluoran, 3-(N-ethyl-N-ethoxypropylamino)-6-methyl-7-anilinofluoran, 3-cyclohexylamino-6-chlorofluoran, 2-methyl-6-p-(p-dimethylaminophenyl)aminoanilinofluoran, 2-methoxy-6-p-(p-dimethylaminophenyl)aminoanilinofluoran, 2-chloro-3-methyl-6-p-(p-phenylaminophenyl)aminoanilinofluoran, 2-diethylamino-6-p-(p-dimethylaminophenyl)aminoanilinofluoran, 2-phenyl-6-methyl-6-p-(p-phenylaminophenyl)aminoanilinofluoran, 2-benzyl-6-p-(p-phenylaminophenyl)aminoanilinofluoran, 3-methyl-6-p-(p-dimethylaminophenyl)aminoanilinofluoran, 3-diethylamino-6-p-(p-diethylaminophenyl)aminoanilinofluoran, 3-diethylamino-6-p-(p-dibutylaminophenyl)aminoanilinofluoran, 2,4-dimethyl-6-[(4-dimethylamino)anilino] fluoran, 3,6,6'-tris(dimethylamino)spiro[fluorene-9,3'-phthalide], 3,6,6'-tris(diethylamino)spiro[fluorene-9,3'-phthalide], 3,3-bis(p-dimethylaminophenyl)-6-

dimethylaminophthalide, 3,3-bis(p-dimethylaminophenyl)phthalide, 3,3-bis-[2-(p-dimethylaminophenyl)-2-(p-methoxyphenyl)ethenyl-4,5,6,7-tetrabromophthalide, 3,3-bis-[2-(p-dimethylaminophenyl)-2-(p-methoxyphenyl)ethenyl-4,5,6,7-tetrachlorophthalide, 3,3-bis[1,1-bis(4-pyrrolidinophenyl)ethylene-2-yl]-4,5,6,7-tetrabromophthalide, 3,3-bis-[1-(4-methoxyphenyl)-1-(4-pyrridinophenyl)ethylene-2-yl]-4,5,6,7-tetrachlorophthalide, 3-(4-diethylamino-2-ethoxyphenyl)-3-(1-ethyl-2-methylindole-3-yl)-4-azaphthalide, 3-(4-diethylamino-2-ethoxyphenyl)-3-(1-octyl-2-methylindole-3-yl)-4-azaphthalide, 3-(4-cyclohexylethylamino-2-methoxyphenyl)-3-(1-ethyl-2-methylindole-3-yl)-4-azaphthalide, 3,3-bis(1-ethyl-2-methylindole-3-yl) phthalide, 3,3-bis(1-octyl-2-methylindole-3-yl) phthalide, mixture of 2-phenyl-4-(4-diethylaminophenyl)-4-(4-methoxyphenyl)-6-methyl-7-dimethylamino-3,1-benzoxazine and 2-phenyl-4-(4-diethylaminophenyl)-4-(4-methoxyphenyl)-8-methyl-7-dimethylamino-3,1-benzoxazine, 4,4'-[1-methylethylidene]bis(4,1-phenyleneoxy-4,2-quinazolinediyl)]bis[N,N-diethylbenzenamine], bis(N-methyldiphenylamine)-4-yl-(N-butylcarbazole)-3-yl-methane.

Especially preferred are fluoran compound such as 3-(4-diethylamino-2-ethoxyphenyl)-3-(1-ethyl-2-methylindole-3-yl)-4-azaphthalide, 3-(4-diethylamino-2-ethoxyphenyl)-3-(1-octyl-2-methylindole-3-yl)-4-azaphthalide, 3-(4-cyclohexylethylamino-2-methoxyphenyl)-3-(1-ethyl-2-methylindole-3-yl)-4-azaphthalide, 3,3-bis(1-ethyl-2-methylindole-3-yl) phthalide, 3,3-bis(1-octyl-2-methylindole-3-yl) phthalide, mixture of 2-phenyl-4-(4-diethylaminophenyl)-4-(4-methoxyphenyl)-6-methyl-7-dimethylamino-3,1-benzoxazine and 2-phenyl-4-(4-diethylaminophenyl)-4-(4-methoxyphenyl)-8-methyl-7-dimethylamino-3,1-benzoxazine, 4,4'-[1-methylethylidene]bis(4,1-phenyleneoxy-4,2-quinazolinediyl)]bis[N,N-diethylbenzenamine], bis(N-methyldiphenylamine)-4-yl-(N-butylcarbazole)-3-yl-methane, 3-methyl-6-p-(p-dimethylaminophenyl)aminoanilino-fluoran, 3-diethylamino-6-p-(p-diethylamino-phenyl)aminoanilino-fluoran, 3-diethylamino-6-p-(p-dibutylaminophenyl)aminoanilino-fluoran, 2,4-dimethyl-6-[(4-dimethylamino)anilino] fluoran, 3,6,6'-tris(dimethylamino)spiro[fluorene-9,3'-phthalide], 3,6,6'-tris(diethylamino)spiro[fluorene-9,3'-phthalide], 3,3-bis(p-dimethylaminophenyl)-6-dimethylaminophthalide, 3,3-bis(p-dimethylaminophenyl)phthalide, 2-diethylamino-6-p-(p-dimethylaminophenyl)aminoanilino-fluoran, 3-dibutylamino-6-methyl-7-(N-formylmethylamino)-fluoran, 2-methyl-6-p-(p-dimethylaminophenyl)aminoanilino-fluoran, 2-methoxy-6-p-(p-dimethylaminophenyl)aminoanilino-fluoran, 3-(N-ethyl-N-ethoxypropylamino)-6-methyl-7-anilino-fluoran, 3-(N-ethyl-N-tetrahydrofurfurylamino)-6-methyl-7-anilino-fluoran, 3-(N-ethyl-N-isobutylamino)-6-methyl-7-anilino-fluoran, 3-(N-methyl-N-propylamino)-6-methyl-7-

anilinofluoran, 3-(N-methyl-N-cyclohexylamino)-6-methyl-7-anilinofluoran, 3-(N-ethyl-N-cyclohexylamino)-6-methyl-7-anilinofluoran, 3-(N-ethyl-p-toluidino)-6-methyl-7-anilinofluoran, 3-(N-ethyl-N-isoamylamino)-6-methyl-7-anilinofluoran, 3-dipentylamino-6-methyl-7-anilinofluoran, 3-dibutylamino-6-methyl-7-anilinofluoran, 3-diethylamino-7-methylfluoran, 3-diethylamino-7-t-butylfluoran, 3-diethylamino-7-carboxyethylfluoran, 3-diethylamino-7-(dibenzylamino) fluoran, 3-dibutylamino-7-dibenzylaminofluoran, 3-diethylamino-6-methylfluoran, 3-diethylamino-6-methyl-7-(4-n-octylanilino) fluoran, 3-diethylamino-7-(4-n-octylanilino) fluoran, 3-diethylamino-7-(4-n-octylamino) fluoran.

All of the above colour forming compounds can be used singly or as a mixture with other colour forming compounds.

The polymeric material usually contains preferably 0,001 to 10% by weight, most preferably 0,01 to 5% by weight of the latent acid. The polymeric material can contain mixtures of two or more of the latent acids.

The amount of colour former in the polymeric material usually is in the range of about 0,001 to 10% by weight, most preferably 0,01 to 5% by weight of the colour former with respect to the total weight of the polymeric material. The polymeric material can contain mixtures of two or more colour formers.

The ratio of latent acid to colour former can e.g. be in the range of 0.1 to 10 parts of colour former per part of latent acid. It is preferred that this weight ratio is about 0.1 to 1.

The polymeric material, the colour former and the latent acid usually form a homogenous mixture. For specific applications, however, compositions can be made in which the latent acid and the colour former are enriched in a part of the polymeric material, e.g. in the surface areas.

The methods for incorporating the colour former and the latent acid into the polymeric material are in principle known. It is e. g. possible, to dissolve the components in a solvent and then to remove the solvent by evaporation. Another possibility is to melt polymeric material together with the colour former and the latent acid to get a homogeneous mixture or to thoroughly knead a mixture of polymeric material, colour former and latent acid.

In another embodiment of this invention, the latent acid (1) preferably is grafted on the polymer material by means known in the art. E.g. the latent acid (1) is converted into a monomer, i.e. by incorporating a functional polymerizable group, or a monomer is used which is functionalized with a latent acid group. This allows a graft polymerization on the existing polymeric material or a copolymerization during the manufacturing the polymeric material.

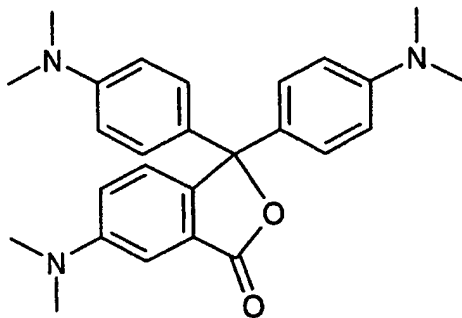
The polymeric material can contain further ingredients, e.g. stabilizers, antioxidants, softeners etc. as are commonly used for polymeric material.

To convert the latent acid into the acid the polymeric material is irradiated. Irradiation in this application especially means irradiation by UV-light and especially by UV-lasers. The lasers used are commercially available. The wavelength of the UV-light preferably is in the range of 285 to 400 nm, more preferably in the range of 285 to 370 nm. The duration of irradiation depends on the components and on the type of UV-source and be easily be determined by simple experiments.

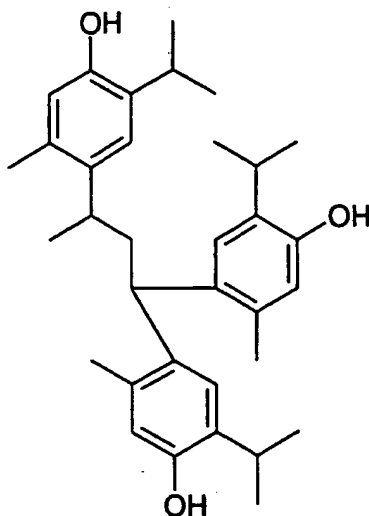
The following non-limitative examples illustrate the invention in more detail. Parts and percentages are by weight, unless otherwise stated.

Examples

Example 1: A polycarbonate film is produced by dissolving polycarbonate together with 1% by weight of the colour former of the formula



and 1% by weight of the latent acid of the formula

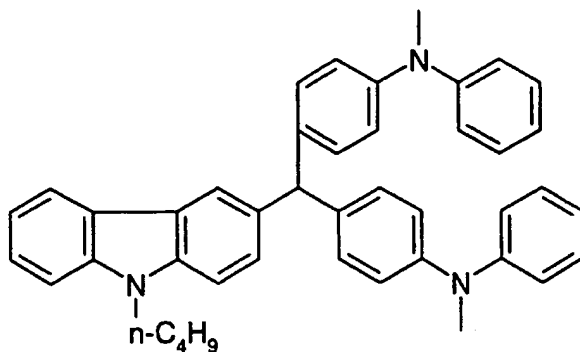


in tetrahydrofurane and then the solvent is evaporated overnight. The polycarbonate film thus produced is irradiated with a UV-laser at 355 nm produces blue marks at the irradiated areas. No background colouration is observed and the lightfastness of the blue marks is good.

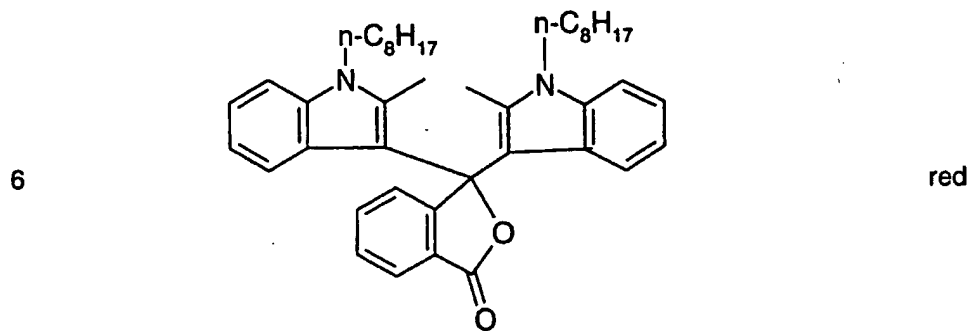
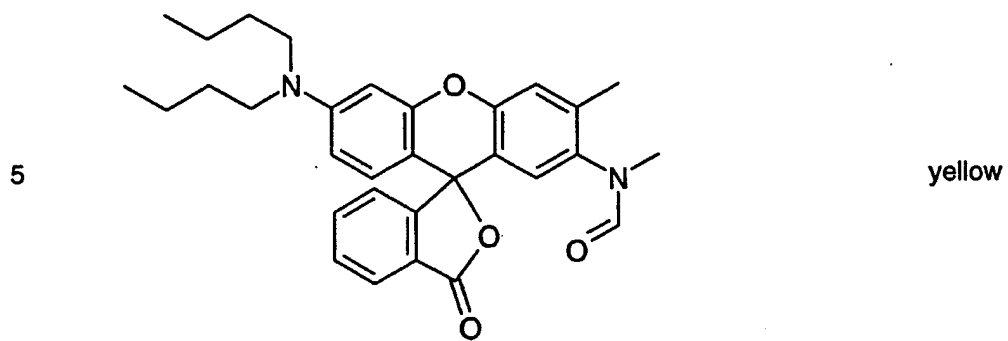
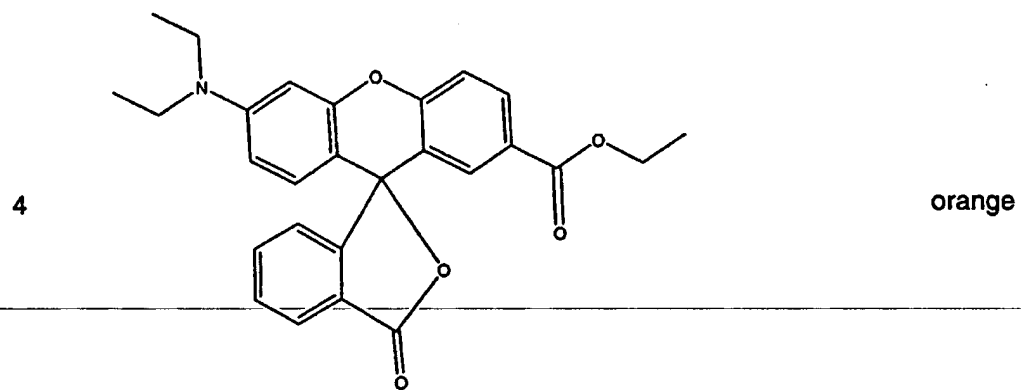
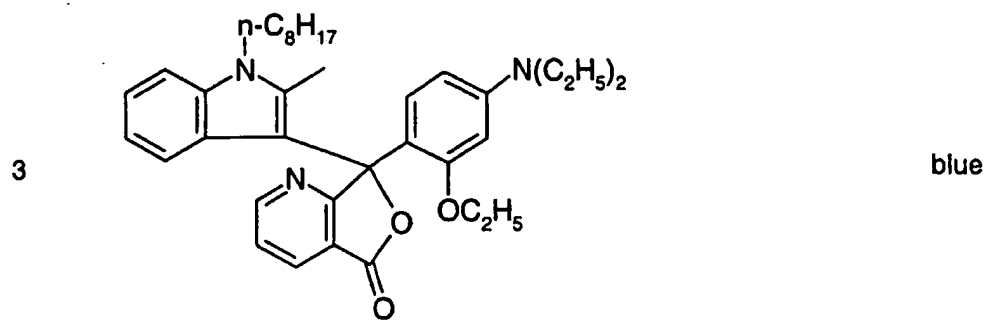
Examples 2 to 8: Repeating example 1 but using 1% of the colour formers of the following formulae gives marks in the given shades having good lightfastness. No background colouration is observed.

Example	Colour former	Shade
---------	---------------	-------

2



blue



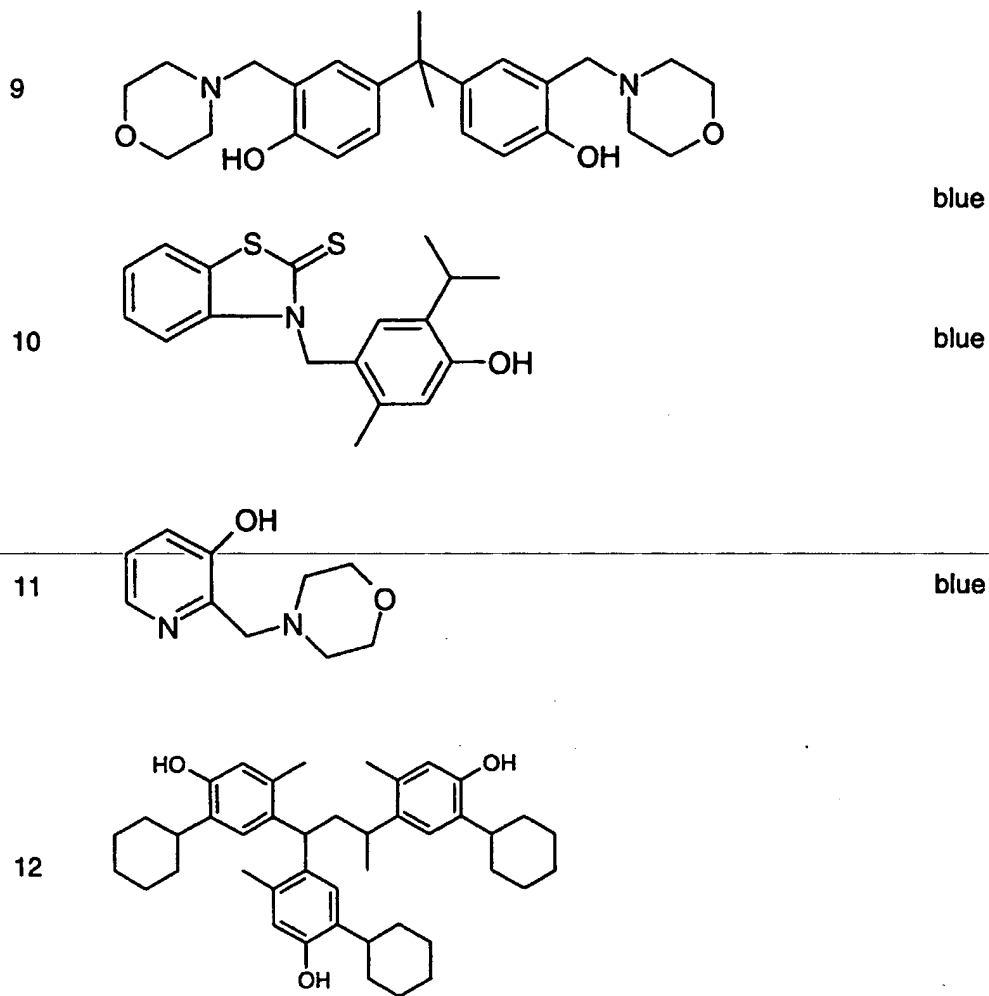
CCN(CC)Cc1ccc2c(c1)oc3ccccc3c2C4(Cc5ccccc5)N(Cc6ccccc6)CC7C(Cc8ccccc8)CCN4C7

8

CCN(CC)c1ccc(cc1)-c2nc3ccccc3n2Oc4ccc(cc4)C(C)(C)c5ccc(cc5)Oc6nc7ccccc7n6-c8ccc(cc8)N(CC)CC

Examples 9 to 12: Repeating example 1 but using 1 % of the latent acids of the following formulae gives marks in the given shades having good lightfastness. No background colouration is observed.

Example Latent Acid Shade



Example 13 to 20: The colour former and latent acid are incorporated in a polycarbonate matrix in the melt form and then extruded and injection moulded to form substantially colourless plaques. When imaged with a UV laser operating at 355 nm clear marks in the given shades with good lightfastness are observed. No colouration of the background is observed.

Example	Colour former (%)	Latent Acid (%)	Remarks
13	1 (0.1)	12 (0.1)	Intense blue mark
14	1 (0.01)	12 (0.01)	Feint Blue mark
15	1 (0.001)	12 (0.001)	Very feint blue mark
16	1 (1)	2 (2)	Intense blue mark
17	1 (2)	2 (1)	Blue mark
18	1(0.5) +2 (0.5)	2(1)	Intense grey-blue mark
19	5 (1)	2 (1)	Intense orange-brown mark
20	4 (1)	2 (1)	Intense orange-brown mark

For examples 13-20 both single pulse marks, text and block images can be created.

Examples 21 to 24: The colour former (1) and latent acid (12) are incorporated in the given matrices by melt processing and are then further processed by conventional means to give the substrates specified below. When imaged with a UV laser operating at 355 nm clear marks in the given shades with good lightfastness are observed. No colouration of the background is observed.

Example	Substrate	%Colour Former (1)	% Latent Acid (12)	Remarks
21	LDPE plaques	0.1	0.1	Intense blue image
22	PP plaques	0.1	0.1	Intense blue image
23	PET cast films	0.1	0.1	Blue image
24	LLDPE blow films	0.1	0.1	Intense blue image

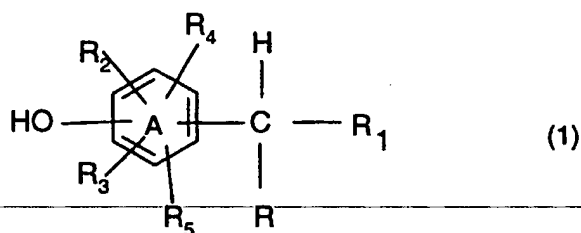
Claims

1. Method of coloring a polymeric material, characterized in that a polymeric material containing

- a latent acid,
- a colour former and
- optionally further ingredients

is irradiated with UV-light.

2. Method according to claim 1, wherein the latent acid is of formula



wherein ring A can contain one or more hetero atoms and/or can contain an anelated ring,

R₁ is hydrogen, alkyl, alkenyl, aryl,

R₂, R₃, R₄ and R₅ independently of each other are hydrogen or a functional substituent, and

R stands for C₁-C₆alkyl, -Z₁-Q₁, or -Z₂-Q₂,

wherein Z₁ is a single bond, S, NH or O, and Q₁ is a heterocyclic ring system having from 5 to 9 ring atoms selected from C, S, O and N, with at least 2 carbon atoms in the ring system, preferably Q₁ stands for morpholine, pyridine, which may be substituted one to three times with C₁-C₄alkyl or hydroxy, mercaptobenzoxazole, mercaptobenzthiazole,

and wherein Z₂ stands for C₁-C₄alkylene, which can be substituted by C₁-C₄alkyl or Q₃,

wherein Q₃ stands for phenyl which can be substituted one to three times with C₁-C₄alkyl,

hydroxy, C₅-C₈cycloalkyl and/or a heterocyclic ring system having from 5 to 9 ring atoms selected from C, S, O and N, with at least 2 carbon atoms in the ring system, and Q₂ stands

for phenyl which can be substituted one to three times with C₁-C₄alkyl, hydroxy, C₅-C₈cycloalkyl and/or a heterocyclic ring system having from 5 to 9 ring atoms selected from C, S, O and N, with at least 2 carbon atoms in the ring system, with the proviso that the hydrogen atom at the C-atom in α-position to R can be split off by irradiation.

3. Method according to any one of claims 1 to 2, wherein R₁ is hydrogen or methyl.

4. Method according to any one of claims 1 to 3, wherein R_2 and R_3 are independently of each other C_1 - C_8 -alkyl and R_4 and R_5 are each hydrogen.
5. Method according to any one of claims 1 to 4, wherein R is a heterocyclic residue, which is bond to the CHR_1 -group via a nitrogen, oxygen or sulfur atom or is a C_1 - C_6 -alkyl, which is unsubstituted or substituted.
6. Method according to any one of claims 1 to 5, wherein R is a radical of mercaptobenzoxazol or mercaptobenzthiazol or C_1 - C_4 -alkyl which is unsubstituted or substituted by unsubstituted phenyl or phenyl carrying 1 to 4 substituents selected from the group consisting of C_1 - C_6 -alkyl, C_1 - C_4 -alkoxy and hydroxy.
7. Method according to any one of claims 1 to 6, wherein the colour former is a triphenylmethane, lactone, benzoxazine, spiropyran or preferably fluoran or phthalide.
8. Method according to any one of claims 1 to 7, wherein the polymeric material contains 0,001 to 10 % by weight, most preferably 0,01 to 5 % by weight of the latent acid.
9. Method according to any one of claims 1 to 8, wherein the polymeric material contains 0.001 to 10 % by weight, preferably 0.01 to 5 % by weight of the colour former with respect to the total weight of the polymeric material.
10. Method according to any one of claims 1 to 9, wherein the polymeric material is irradiated with a UV-laser using UV-light of 285 to 400 nm.

INTERNATIONAL SEARCH REPORT

International Application No

PCT/EP 02/06105

A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 G03C1/73 B41M5/00 G03F7/105 G03C1/00 G03F7/00
 B41M5/26 B41M5/30 C09B67/00

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 C08K B41M G03C G03F C09B

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the International search (name of data base and, where practical, search terms used)

WPI Data, PAJ; EP0-Internal

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	EP 0 290 750 A (XYTRONIX INC) 17 November 1988 (1988-11-17) page 1, line 10 - line 20; claims 1-5,13,14,30; examples 1-31 page 4, line 5 -page 5, line 22 ---	1,9,10
X	US 4 343 885 A (REARDON JR EDWARD J) 10 August 1982 (1982-08-10) column 2, line 1 -column 25, line 43; claims 1-16,19,30,41,45,46 column 17, line 37 - line 45 --- -/--	1,9,10



Further documents are listed in the continuation of box C.



Patent family members are listed in annex.

* Special categories of cited documents:

A document defining the general state of the art which is not considered to be of particular relevance

E earlier document but published on or after the international filing date

L document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

O document referring to an oral disclosure, use, exhibition or other means

P document published prior to the international filing date but later than the priority date claimed

T later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

X document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

Y document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

Z document member of the same patent family

Date of the actual completion of the international search

16 August 2002

Date of mailing of the international search report

29/08/2002

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International Application No

PCT/EP 02/06105

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	<p>EP 0 600 441 A (DAINIPPON INK & CHEMICALS ;KIRIN BREWERY (JP)) 8 June 1994 (1994-06-08) page 2, line 3-6 page 3, line 14 -page 4, line 16 page 5, line 27 - line 34 page 6, line 33 - line 41 page 7, line 4-10; claims 1,6,10 -----</p>	1,9,10
X	<p>EP 0 720 053 A (TOKYO OHKA KOGYO CO LTD) 3 July 1996 (1996-07-03) page 2, line 54 - line 58 page 3, line 5 - line 41; claims 1,3-8 page 7, line 8 -page 14, line 4 page 14, line 31 - line 34 -----</p>	1,9,10
X	<p>US 5 677 107 A (NECKERS DOUGLAS C) 14 October 1997 (1997-10-14) column 4, line 53 -column 5, line 29 column 9, line 56 -column 10, line 11; claims 1,4,19,20,22 -----</p>	1,9,10

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Information on patent family members

International Application No

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